

An anionic binuclear complex of tungsten(II), $[(\mu\text{-Cl})_3\{\text{W}(\text{SnCl}_3)(\text{CO})_3\}_2]^-$, and its reactivity towards norbornene.

Autorzy

Anna Malinowska

Andrzej Kochel

Teresa Szymańska-Buzar

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Streszczenie

An anionic binuclear complex of tungsten(II), $[(\mu\text{-Cl})_3\{\text{W}(\text{SnCl}_3)(\text{CO})_3\}_2]^-$ (**1**⁻), containing the protonated piperidine molecule $[\text{Hpip}]^+$ as the counter ion, has been obtained during crystallization of the product from reaction between $[\text{W}(\text{CO})_4(\text{pip})_2]$ and SnCl_4 in dichloromethane solution, and its molecular structure has been elucidated by single-crystal X-ray diffraction studies. The chemical properties of complex **1** were investigated by IR and NMR spectroscopy in solution and its catalytic activity was checked in reaction with norbornene (NBE). In the presence of complex **1**, NBE transformed to a new olefin, 2,2'-binorbornylidene with ca. 50% yield in dichloromethane solution. The spectroscopic characteristics of complex **1**⁻ were compared with those of the reinvestigated analogue compound $[(\mu\text{-Cl})_3\text{W}_2(\text{SnCl}_3)(\text{CO})_7]$ (**2**). The ¹¹⁹Sn and ¹³C NMR data indicated that in dichloromethane solution complex **2** transformed to the ionic complex **1**⁻.

Słowa kluczowe

Tungsten(II), Stannyl ligand, Heterobimetallic complex, X-ray crystal structure, Catalytic activity, Olefin dimerization, Carbene formation

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